AMENDMENTS TO THE CLAIMS

- 1. (Original) A kit for activating gene transfer, said kit comprising a gene transfer activating compound, packaged in a suitable container together with instructions for use to activate gene transfer.
- 2. (Original) The kit of claim 1 wherein said gene transfer activating compound has a molecular weight of between 300 and 2000.
- 3. (Original) The kit of claim 1 wherein said gene transfer compound is selected from the group consisting of:

wherein Q is nitrogen or oxygen, wherein each occurrence of R^1 independently is H, CH_3 , CH_2CH_3 or a nullity, wherein R^2 is C_1 - C_{18} alleyl, C_2 - C_{18} ether, C_2 - C_{18} thioether, C_2 - C_{18} secondary or tertiary amine,

wherein A is

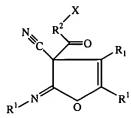
wherein R^3 is H, C_1 – C_6 alkyl, or a heteroatom substituted C_1 – C_6 alkyl where the heteroatom is oxygen, nitrogen, or sulfur, wherein R^4 is C_2 – C_6 amide, or =N– R^5 where R^5 is C_7 – C_{12} aryloxyl, C_1 – C_6 hydronyl, carbonyl, carboxyl, or acyl, imidazyl, pyrazyl, thiazyl, or oxazyl, wherein X is H, F, Cl or Br, wherein Z is oxygen or sulfur.

- 4. (Original) The kit of claim 1 wherein said gene transfer compound is bouvardin.
- 5. (Original) The kit of claim 3 wherein said gene transfer compound is that of

6. (Original) The kit of claim 3 wherein said gene transfer compound is that of

structure I, wherein A and each occurrence of Q together are

- 7. (Original) The kit of claim 3 wherein said gene transfer compound is that of structure II wherein Q is nitrogen and R^2 is C_1-C_{18} alkyl.
 - 8. (Original) The kit of claim 7 wherein R^4 is $=N-R^5$.
 - 9. (Original) The kit of claim 7 wherein X is Cl or Br.
 - 10. (Original) The kit of claim 3 wherein said gene transfer compound is that of



structure III wherein Q in each occurrence together are R1/

- 11. (Original) The kit of claim 10 wherein said gene transfer compound is that of structure II or VII wherein each occurrence of R¹ is H, or CH₃.
- 12. (Original) The kit of claim 3 wherein said gene transfer compound is that of structure V wherein Q in each occurrence is oxygen.

13. (Original) The kit of claim 3 wherein said gene transfer compound is that of structure VI wherein Q in each occurrence is oxygen.

$$Q \longrightarrow R^1$$

- 14. (Original) The kit of claim 13 wherein A is
- 15. (Original) The kit of claim 3 wherein said gene transfer compound is that of structure VII wherein Q in each non-aromatic substituent occurrence is oxygen.
 - 16. (Original) The kit of claim 15 wherein R¹ in each occurrence is H.
- 17. (Original) The kit of claim 3 wherein said compound is selected from the group consisting of: NSC73609, NSC82090, NSC101492, NSC102821, NSC106191, NSC108613, NSC109325, NSC128720, NSC143491, NSC259968, NSC373989 and NSC675865.

Claims 18-27 (Canceled)

28. (Currently Amended) The process of claim 27 A process for activating gene transfer of a vector to a cell comprising the steps of:

contacting a cell with a recombinant gene transfer vector; and

administering a gene transfer activating compound to the cell, such that transfer of the vector to the cell is activated;

wherein the gene transfer activating compound is selected from the group consisting of:

wherein Q is nitrogen or oxygen, wherein each occurrence of R^1 independently is H, CH_3 , CH_2CH_3 or a nullity, wherein R^2 is C_1 - C_{18} alleyl, C_2 - C_{18} ether, C_2 - C_{18} thioether, C_2 - C_{18} secondary or tertiary amine,

wherein A is

$$\begin{array}{c|c} & & & \\ & & &$$

wherein R^3 is H, C_1 – C_6 alkyl, or a heteroatom substituted C_1 – C_6 alkyl where the heteroatom is oxygen, nitrogen, or sulfur, wherein R^4 is C_2 – C_6 amide, or =N– R^5 where R^5 is C_7 – C_{12} aryloxyl, C_1 – C_6 hydronyl, carbonyl, carboxyl, or acyl, imidazyl, pyrazyl, thiazyl, or oxazyl, wherein X is H, F, Cl or Br, wherein Z is oxygen or sulfur.

29. (Currently Amended) The process of claim 27 A process for activating gene transfer of a vector to a cell comprising the steps of:

contacting a cell with a recombinant gene transfer vector; and

administering a gene transfer activating compound to the cell, such that transfer of the vector to the cell is activated;

wherein the gene transfer activating compound is selected from the group consisting of: NSC73609, NSC82090, NSC101492, NSC102821, NSC106191, NSC108613, NSC109325, NSC128720, NSC143491, NSC259968, NSC373989 and NSC675865.

Claims 30-35 (Canceled)

36. (Original) A process for determining the efficacy of a putative gene transfer activating compound to activate gene transfer, comprising the steps of:

administering a test compound to a first cell;

contacting the first cell with a first amount of a recombinant vector;

contacting a second cell with a second amount of the recombinant vector, the second amount of the recombinant vector substantially equal to the first amount;

measuring a gene transfer indicator in the first cell to obtain a test measurement;

measuring the gene transfer indicator in the second cell to obtain a control measurement; and

comparing the test measurement and the control measurement to determine the efficacy of the putative gene transfer activating compound to activate gene transfer.

- 37. (Original) Use of a compound of Formulae I-VII for use as a gene transfer activating compound.
- 38. (Original) The use of claim 37 wherein said gene transfer activating compound has a molecular weight of between 300 and 2,000.
- 39. (Original) The use of claim 37 wherein said gene transfer compound is selected from the group consisting of:

wherein Q is nitrogen or oxygen, wherein each occurrence of R^1 independently is H, CH_3 , CH_2CH_3 or a nullity, wherein R^2 is C_1 - C_{18} allyl, C_2 - C_{18} ether, C_2 - C_{18} thioether, C_2 - C_{18} secondary or tertiary amine,

wherein A is

wherein R^3 is H, C_1 – C_6 alkyl, or a heteroatom substituted C_1 – C_6 alkyl where the heteroatom is oxygen, nitrogen, or sulfur, wherein R^4 is C_2 – C_6 amide, or =N– R^5 where R^5 is C_7 – C_{12} aryloxyl, C_1 – C_6 hydronyl, carbonyl, carboxyl, or acyl, imidazyl, pyrazyl, thiazyl, or oxazyl, wherein X is H, F, Cl or Br, wherein Z is oxygen or sulfur.

- 40. (Original) The use of claim 37 wherein said gene transfer compound is bouvardin.
- 41. (Original) The use of claim 39 wherein said gene transfer compound is that of structure I, wherein A is , and Q is nitrogen in each occurrence.
 - 42. (Original) The use of claim 39 wherein said gene transfer compound is that of

structure I, wherein A and each occurrence of Q together are
$$N$$

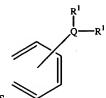
- 43. (Original) The use of claim 39 wherein said gene transfer compound is that of structure II wherein Q is nitrogen and R^2 is C_1 – C_{18} alkyl.
 - 44. (Original) The use of claim 43 wherein R^4 is $=N-R^5$.

- 45. (Original) The use of claim 43 wherein X is Cl or Br.
- 46. (Original) The use of claim 39 wherein said gene transfer compound is that of

$$\begin{array}{c|c}
N & X \\
R^2 & O \\
R^1 & N & O
\end{array}$$

structure III wherein Q in each occurrence together are R1/

- 47. (Original) The use of claim 46 wherein said gene transfer compound is that of structure II or VII wherein each occurrence of R¹ is H, or CH₃.
- 48. (Original) The use of claim 39 wherein said gene transfer compound is that of structure V wherein Q in each occurrence is oxygen.
- 49. (Original) The use of claim 39 wherein said gene transfer compound is that of structure VI wherein Q in each occurrence is oxygen.



- 50. (Original) The use of claim 49 wherein A is
- 51. (Original) The use of claim 39 wherein said gene transfer compound is that of structure VII wherein Q in each non-aromatic substituent occurrence is oxygen.

- 52. (Original) The use of claim 51 wherein R¹ in each occurrence is H.
- 53. (Original) The use of claim 39 wherein said compound is selected from the group consisting of: NSC73609, NSC82090, NSC101492, NSC102821, NSC106191, NSC108613, NSC109325, NSC128720, NSC143491, NSC259968, NSC373989 and NSC675865.
 - 54. (Canceled)